

(12) UK Patent Application (19) GB (11) 2 114 566 A

(21) Application No 8303280
(22) Date of filing 7 Feb 1983
(30) Priority data
(31) 8203636
8226006

(32) 9 Feb 1982
13 Sep 1982
(33) United Kingdom (GB)
(43) Application published
24 Aug 1983

(51) INT CL³
C07D 333/36 261/14
307/66
(52) Domestic classification
C2C 1510 1600 200 202
215 220 226 22Y 246 247
1340 1341 1370 1371 1380
1382 1386 1400 1410 1414
1432 1440 1450 1470 1472
1492 1510 254 255 256
25Y 27X 281 282 28X 30Y
247 250 251 252 253 254
30Y 311 313 314 31Y 321
322 323 326 327 32Y 332
364 365 366 367 368 36Y
371 373 37Y 396 397 43X
440 442 446 450 453 45X
45Y 470 471 490 491 575
579 584 595 59Y 601 337
339 340 342 346 34Y 351
352 354 355 35Y 360 361
364 652 658 65X 670 671
672 675 677 678 67X 680
695 72Y 743 758 776 805
80Y AA 601 602 610 614
620 621 623 624 625 627
628 62X 634 635 638 63X
644 652 AA BL KA KF KJ
KK KM KP KQ KR KS KV
KY KZ LH LL LS LT MF ML
NG RE RM

U1S 1306 1347 C2C
(56) Documents cited
GB 1497536

(58) Field of search
C2C

(71) Applicants
Sandoz Ltd
(Switzerland),
35 Lichtstrasse,
CH-4002 Basle,
Switzerland

(72) Inventors
Karl Seckinger
Fred Kuhnen
Karlheinz Miltner

(74) Agent and/or
Address for Service
B. A. Yorke and Co.,
98 The Centre,
Feltham,
Middlesex TW13 4EP

(54) Herbicidal chloroacetamides

(57) Novel 5-membered
heteroaromatic compounds of formula I
$$\text{ArN(Y)COCH}_2\text{Cl}$$

wherein Ar is a 5-membered
heteroaromatic group
comprising 1 or 2
heteroatoms selected from
O, S and N and linked by a
ring C-atom to the N-atom
of the N(Y)COCH₂Cl group
to which it is bound,
whereby where Ar is
pyrazolyl, said
N(Y)COCH₂Cl group is in
the 4-position,

and Y is as specified in the description,
are useful as herbicides. Novel starting
materials and intermediates are also
described.

GB 2 114 566 A

SPECIFICATION

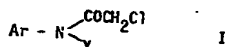
Chloroacetamides

- 5 The present invention relates to novel 5-membered heteroaromatic compounds bearing at one ring C-atom a N-substituted chloroacetyl amino group, their use as herbicides, agricultural compositions for facilitating such use and the preparation of the novel compounds of the invention.

Various herbicidal N-substituted α -halogenacetanilides are known.

- The U.S. Patent Spec. 4,282,028 discloses N-substituted - N - 2,5 - dialkyl - pyrrol - 1 - yl) haloacetamides having herbicidal and plant - growth - regulating activity. The need exists for still more effective herbicides. The novel 5-membered heteroaromatic compounds are particularly effective herbicides having an appropriate soil persistence.

- 20 The present invention provides compounds of formula I



- wherein Ar is a 5-membered heteroaromatic group comprising 1 or 2 ring heteroatoms selected from O, S and N, and linked by a C-ring atom to the N-atom of the N(Y)COCH₂Cl group to which it is bound, whereby, where Ar is pyrazolyl, said N(Y)COCH₂Cl group is in the 4-position,

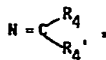
- and Y is allene, CH₂-CH=C=CH₂, or is a hydrocarbon selected from C₁₋₈alkyl, C₃₋₈alkenyl, C₃₋₈alkinyl, C₃₋₈cycloalkyl, C₅₋₈cycloalkenyl, C₃₋₈cycloalkyl - C₁₋₅alkyl, whereby such hydrocarbon is unsubstituted or substituted by halogen selected from F, Cl or Br; or is a group CH(R₁)-COY₁,

- wherein R₁ is H or C₁₋₅alkyl and Y₁ forms together with the CO-group to which it is bound an ester or amide function;

- or is a group R₂-A₂, wherein R₂ is CH₂ or CH₂-CH₂ unsubstituted or substituted by C₁₋₅alkyl and

- 40 Az is an aromatic heteroring selected from a di- or triazole linked by one of its nitrogen atoms to R₂, a 5-membered heteroring linked by a C-atom of said ring to R₂ and having 1 to 3 heteroatoms selected from the group consisting of O, S or N, and a pyrimidine group, or is a 2 - oxo - 1 - pyrrolidinyl group in which one CH₂ group can be replaced by O, S or NCH₃ and 5-oxo- and/or bicyclic benz[c] fused derivatives of such 2 - oxo - 1 - pyrrolidinyl group or is a group of A-O-R₃,

- 50 wherein R₃ is H, hydrocarbon selected from the group consisting of C₁₋₈alkyl, C₃₋₈alkenyl, C₃₋₈alkinyl, C₃₋₈cycloalkyl, C₅₋₈cycloalkenyl or C₃₋₈cycloalkyl-C₁₋₅alkyl, whereby the hydrocarbon is unsubstituted or substituted; or a group



- 55 wherein R₄ is hydrocarbon selected from C₁₋₅alkyl, C₃₋₅alkenyl, C₃₋₅alkinyl, C₃₋₅cycloalkyl, C₅₋₈cycloalkenyl, C₃₋₈cycloalkyl-C₁₋₅alkyl, whereby the hydrocarbon is unsubstituted or substituted by

halogen selected from F, Cl or Br; or is allene

- 60 R₄' is H or has one of the meanings defined for R₄ and A is a hydrocarbon moiety, which may be linked with R₃ to form a saturated oxygen containing heterocyclic ring comprising 1 or 2 oxygens as heteroatom, and whereby the N and O atoms to which it is bound are separated by up to 3 C-atoms; or is a group CH - C = NOR₄,



- wherein R₄ is as defined above, R₅ and R₅' are independently H or CH₃ or R₅ together with R₅' are (CH₂)₃ or (CH₂)₄

- 70 or is a group CH-N(CH₃)COCH₃



wherein R₆ is H or C₁₋₃alkyl.

The Ar group may be unsubstituted or substituted.

- 75 Where Ar is substituted it may bear substituents in any possible position; preferred positions of such substituents are in o-position, particularly in o,o'-position of the chloroacetamide group, whereby additional substituents may be present. Examples of suitable substituents of Ar are halogen selected from F, Cl and Br; C₁₋₄alkyl unsubstituted or substituted by halogen (F, Cl, Br), C₁₋₄alkoxy or C₃₋₆cycloalkyl; C₂₋₄alkenyl unsubstituted or substituted by C₁₋₄alkoxy; C₃₋₆cycloalkyl; formyl or C₂₋₄alkanoyl and functional derivatives such as oximes, acetals and ketals thereof (e.g. C(=NOC₁₋₄alkyl) - C₁₋₃alkyl, C(O-C₁₋₄alkyl)₂ - C₁₋₃alkyl, CH(O-C₁₋₄alkyl)₂ etc.); C₁₋₄alkyl-S, C₁₋₄alkyl-SO, C₁₋₄alkyl-SO₂; C₁₋₅alkoxycarbonyl; C₁₋₄alkoxy unsubstituted or substituted by halogen or C₁₋₄alkoxy; C₂₋₄alkenyloxy, C₂₋₄alkinyloxy; hydroxy and hydroxymethyl and esters thereof (e.g. esters with an organic carboxylic acid, for example formic acid, a C₂₋₅alkane carboxylic acid or an halogenated derivative thereof, such as acetic acid or chloroacetic acid).

- Any optional substituent(s) of Ar which is not in o-position of the N-substituted chloroacetyl amino group is preferably selected from C₁₋₄alkyl (e.g. CH₃), halogen (e.g. Cl, Br) and C₁₋₄alkoxycarbonyl (e.g. COOCH₃).

- Where Ar contains 2 ring heteroatoms, one of them is N; the other is preferably selected from O or S.

- A preferred sub-group of compounds of formula I, are compounds wherein Ar contains either one ring heteroatom (pyrrolyl, thienyl, furanyl) or 2 adjacent ring heteroatoms (isoxazolyl, isothiazolyl, pyrazolyl). The N-substituted chloroacetyl amino group is preferably tied to the ring C-atom in β -position of the ring heteroatom(s); thus in the 3-position where Ar is pyrrol, thiophene or furan and in the 4-position where Ar is isoxazole, isothiazole or pyrazole. A preferred sub-group of the latter group are those compounds wherein the Ar-group is substituted in ortho-, more preferably in ortho - ortho' - position of the N-substituted acetyl amino group, especially when the substituent(s) are selected from the group specified hereinbefore. Any ring N-atom may be

Certain of the Chemical formula(e) appearing in the printed specification were submitted after the date of filing, the formula(e) originally submitted being incapable of being satisfactorily reproduced.

unsubstituted or substituted; where it is substituted, it is preferably substituted by C₁₋₄alkyl, especially CH₃ or C₂H₅. Ar signifies preferably a thiophene, isothiazole and isoxazole, more preferably

- 5 isothiazole or thiophene ring. A particularly preferred sub-group of compounds of formula I, are compounds wherein Ar is thiophene, particularly 3-thienyl which is at least 2,4-disubstituted, especially those 3-thienyl compounds wherein the substituents in 2- and 4-position selected from C₁₋₄alkyl and C₁₋₄alkoxy.

Where Y, R₃, R₄ and R₄' are hydrocarbon substituted by halogen, such halogen is preferably Cl or Br; any such substituted hydrocarbon is particularly 15 monosubstituted. Where Y is halogen substituted hydrocarbon, the halogen is in general preferably not in the 1-position where the use as herbicide is intended (such 1-halogenated compounds are in general unstable; they are however useful as valuable intermediate for the production of other compounds of formula I, e.g. for those wherein Y is alkoxyalkyl or pyrazolylmethyl); a suitable example of such significance of Y is 2-Cl-ethyl.

Where Y, R₃, R₄ and/or R₄' are C₁₋₆alkyl, C₃₋₆alkenyl or C₃₋₆alkinyl, they preferably have up to 5 C-atoms. Where Y, R₃, R₄ and/or R₄' are or contain cycloalkyl or cycloalkenyl such cyclic hydrocarbon groups contain preferably up to 6 C-atoms.

Where R₁ is C₁₋₅alkyl it is preferably CH₃ or C₂H₅, particularly CH₃. The term ester or amide function used in connection with the meaning of Y₁ is intended to embrace any function obtainable by the reaction of the COOH group of an acid with any organic compound reacting with such acid with the elimination of water e.g. an alcohol, an amine, a mercaptan, an oxime, a hydrazine, hydrazide or hydrazone. Examples of suitable ester significances of Y₁ are C₁₋₅alkoxy-CO, C₃₋₅alkinylO-CO etc. Examples of suitable amide significances of Y₁ are C₁₋₄alkylamino-CO, di(C₁₋₄alkyl) amino-CO, CO-NHNH₂ and CO-NH-N=C(C₁₋₃alkyl)₂.

Where R₂ is CH₂ it may be substituted by 1 or 2, preferably 1 C₁₋₅alkyl. Where R₂ is CH₂CH₂ it may be substituted by up to 4 C₁₋₅alkyl group and is preferably mono- or disubstituted. Where CH₂CH₂ is disubstituted, the substituents are preferably at different C-atoms.

Preferred C₁₋₅alkyl substituents of R₂ are CH₃ and C₂H₅, particularly CH₃. Where R₂ is substituted by C₁₋₅alkyl, it is preferably monosubstituted. R₂ is especially CH₂ or CH(CH₃).

Where Az is one of the above defined aromatic heterorings, such ring may be unsubstituted or substituted; suitable substituents of such Az significances are e.g. one or more C₁₋₅alkyl (such as CH₃, C₂H₅) groups; such substituted aromatic heterorings are preferably mono- or di-substituted.

Where Az is di- or triazole it is preferably 1-pyrazolyl, 1-imidazolyl, 1,2,3-triazol-1-yl or 1,2,4-triazole-1-yl, unsubstituted or substituted, particularly 1-pyrazolyl, 3,5-di-CH₃-1-pyrazolyl.

Where Az is the above defined 5-membered heteroring linked by a ring C-atom to R₂, such ring is e.g. a furyl, a thienyl, a thiazolyl, an isothiazolyl, an oxazolyl, an isoxazolyl, a 1,2,4- or 1,3,4-thia- or

oxadiazol or a 1,3,4-triazolyl group, unsubstituted or substituted, e.g. by one or two substituents of the group C₁₋₄alkyl, C₁₋₄alkylthio, C₁₋₄alkoxy; typical examples of such significances are 2-furyl, 2-thienyl, 2-CH₃-4-thiazolyl, 3-CH₃-5-isoxazolyl, 2-C₂H₅-1,3,4-oxadiazol-5-yl, 3-CH₃-1,2,4-oxadiazol-5-yl, 1-CH₃-2-CH₃S-1,3,4-triazol-5-yl and 2-CH₃-1,3,4-triazol-5-yl.

Where Az is a pyrimidine group, this may be linked by its 2-, 4- or 5-C-atom to R₂, and is e.g. 2-pyrimidinyl.

Where Az is the above defined bicyclic 2-oxo-1-pyrrolidinyl derivative, the benzo-moiety may be unsubstituted or substituted, e.g. by halogen; a suitable example of such significance is 2-oxo-1,3-benzothiazol-3-yl and the 4-chloro derivative thereof.

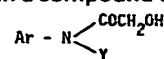
Where R₃ is the above defined hydrocarbon, such hydrocarbon may be unsubstituted or substituted, e.g. by halogen, cyano, an azole group such as 1-pyrazolyl and/or by C₁₋₄alkoxy; such substituted hydrocarbon significances of R₃ are preferable monosubstituted. Examples of such R₃ significances are CH₂CH₂Cl, (1-pyrazolyl)-CH₂, CH₂CH₂CN, 3,5-dimethyl-1-pyrazolyl.

Suitable meanings of A-O-R₃, when A is linked with R₃ to form a ring, are e.g. 1,3-dioxolane-4-yl - C₁₋₃alkylene, 1,3-dioxolane-2-yl - C₁₋₃alkylene and 2- or 3-tetrahydrofuryl - C₁₋₃alkylene.

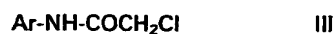
Where A is not linked with R₃ to form a ring, A is preferably a non-aromatic, particularly a C₁₋₈alkylene moiety separating the O- and N-atom to which it is bound by 1 to 3, preferably 1 or 2 C-atoms. A suitable significance of A is a CH₂ or CH₂-CH₂ group or monomethylated derivatives thereof, particularly CH₂, CH(CH₃), CH₂CH₂ and CH(CH₃)CH₂; R₃ is then especially C₁₋₃alkyl, such as CH₃, C₂H₅ and nC₃H₇.

A may be unsubstituted or substituted, e.g. by C₁₋₅alkoxy. Any substituted A significance is preferably monosubstituted.

The present invention also provides processes for producing a compound of formula I comprising a) substituting in a compound of formula II



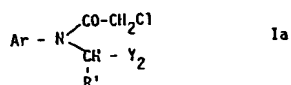
- 110 wherein Ar and Y are as defined above, the HO group of the N-hydroxyacetyl group by Cl, b) reacting a compound of formula III



- 115 wherein Ar is as defined above, with a compound of formula IV

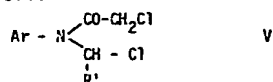


- 120 wherein Y is as defined above, and L is a leaving group capable of being split off under the N-alkylation reaction conditions, c) obtaining a compound of formula Ia



wherein R' is H or C₁₋₃alkyl,
Y₂ is Az', OR₃, N(CH₃)COCH₃ and
Az' is a di- or triazole linked by one of its nitrogen
atoms to CHR'

- 5 and Ar and R₃ are as defined above, by reacting a
compound of formula V



wherein Ar and R' are as defined above, with a
reactive derivative of a compound of formula VI
VI

wherein Y₂ is as defined above,

- 10 d) N-acylation of a compound of formula VII



wherein Ar and Y are as defined above, with
15 chloroacetyl chloride, or a reactive functional deriva-
tive thereof.

Process a) of the invention can be carried out by
conventional manner under conditions known for
the substitution of an OH group by a Cl.

- 20 Such substitution can for example be effected by
treating a compound of formula II with a chlorinating
agent, such as thionyl chloride under conditions
known per se for analogous reactions.

According to a variante of this chlorination pro-
cess, the compounds of formula II are first con-
verted into the corresponding sulphonyloxy deriva-
tives, e.g. by O-sulphonation with the aid of a sulphonyl
halide, and such sulphonyloxy derivatives then
converted into the desired compounds of formula I
30 by nucleophilic substitution of the sulphonyloxy
group by chlorine.

Reactants supplying the Cl anions required for
such nucleophilic substitution are e.g. alkalimetal
chlorides such as NaCl, quaternary tetrabutylammo-
nium chloride or 4 - dimethylaminopyridine - hyd-
rochloride. Such substitution is conveniently carried
out in CH₂Cl₂ or in an aqueous/organic two-phase
system, wherein the organic phase is e.g. a hydro-
carbon such as toluene, in the presence of a suitable
40 phase transfer catalyst, preferably with heating e.g.
at 40° to 120°C.

Process b) may be carried out by conventional
manner under conditions known for the N-alkylation
of amides. The reaction is advantageously carried
out in a solvent which is inert under the reaction
conditions e.g. dimethoxyethane or acetonitrile or in
an aqueous/organic two-phase system in the pre-
sence of a phase transfer catalyst.

Suitable meanings of L (in formula IV) are Cl, Br or
50 the sulphonyloxy moiety of an organic sulphonic
acid such as mesyloxy or p-tosyloxy.

The compounds of formula III are preferably used
in salt form, more preferably in alkalimetal salt form,
e.g. the sodium salt form. Such salts are obtained in
55 conventional manner by reaction of the compound
of formula III with a base such as an alkalimetal
amide, hydride, hydroxide or alcoholate.

For the preparation of compounds of formula Ia
according to process c), the compounds of formula V
are suitably used in the form of the alkalimetal salt,
60 e.g. as sodium salt. Where Y₂ is a di- or triazole, the
compound of formula V may also be used as
N-tri-alkylsilyl (e.g. N-trimethylsilyl) compound. The
reaction can be effected in conventional manner.

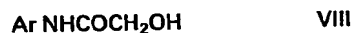
65 The reaction of process d) can likewise be effected
in conventional manner, under conditions known for
the N-chloroacetylation of amines. Where
ClCOCH₂Cl is used as N-chloroacetylation agent
such reaction is conveniently carried out in the
70 presence of an acid binding agent, such as K₂CO₃.

The compounds of formula I may be recovered
from the reaction mixture in which it is formed by
working according to established procedures.

It will be appreciated that interconversion of one
75 compound of the invention to another, e.g. conver-
sion of an acid group to an ester group, of a carbonyl
group to an oxime, of a haloalkyl group to an ether
group (or vice versa) etc., may be carried out in
conventional manner. As will also be appreciated,
80 the compounds of the invention may possess one or
more asymmetric centres and may therefore exist in
optically active, diastereomeric, racemic or geometric
isomer forms. In general such compounds are
employed as mixtures in the herbicide method and
85 compositions of the invention, even though separa-
tion may be effected by known procedures.

The compounds of formulae II, III, V and VII are
novel and are part of the invention.

The compounds of formula II may be obtained by
90 ammonolysis of an ester of the compounds of
formula VIII



95 wherein Ar is as defined above,
with a carboxylic acid, and subsequent introduction
of the group Y (as defined above) in the compounds
of formula VIII by N-alkylation. Such esters are
obtained by acylation of the corresponding com-
pounds of formula IX



wherein Ar is as defined above,
105 with the appropriate ester of HO-CH₂COCl, e.g. with
CH₃CO - OCH₂ - COCl.

The compounds of formula III may also be
obtained by acylation of a compound of formula IX
with chloroacetyl chloride.

110 Compounds of formula V (which are in fact a
sub-group of compounds of formula I) may be
obtained by reaction of a compound of formula IX
with an appropriate aldehyde and reaction of the
Schiff base thus obtained with chloroacetyl chloride.

115 The compounds of formula VII may be obtained by
N-alkylation of a compound of formula IX. Such
alkylation can be carried out in conventional man-
ner, with the corresponding alkylating agents (e.g.
halogenides), or where appropriate, reductively via
120 the Schiff base or amide.

Many of the compounds of formula IX are novel.
A particular group of valuable novel compounds

of formula IX are 3-aminothiophenes substituted in 2- and 4-position by a group selected from C₁₋₄alkyl and C₁₋₄alkoxy (compounds of formula IXa). The compounds of formula IXa also form part of the invention.

The compounds of formula IX may be obtained by reduction of the corresponding NO₂ compounds, e.g. by catalytic hydrogenation under hydrogen pressure in the presence of palladium. Compounds of formula IXa having a CH₃ group in 2- or 4-position may be obtained by reduction of the corresponding thiophene-carboxylate with the aid of a complex hydride such as sodium bis(methoxyethoxy)aluminum hydride. Where appropriate, the compounds of formula IXa may also be obtained from the corresponding carbamate esters, e.g. the benzyl carbamate, by hydrolysis. The carbamates, used as starting material, may for example be obtained starting from the corresponding acids, via their azides followed by a Curtius reaction.

Insofar as the production of starting material is not described herein, these compounds are known, or may be produced and purified in accordance with known processes or in a manner analogous to processes described herein or to known processes.

The compounds of formula I are useful because they control or modify the growth of plants. By plants it is meant germinating seeds, emerging seedlings and established vegetation including underground portions.

In particular, the compounds are useful as herbicides as indicated by i.a. the damage caused to both monocotyledonous and dicotyledonous plants: such as *Lepidium sativum*, *Avena sativa*, *Agrostis alba* and *Lolium perenne* in tests by test dosages equivalent to an application rate of from 1.4 to 5.6 kg/ha after pre- or post-emergence application. In view of their herbicidal effect the compounds of the invention are indicated for use in combatting dicotyledonous and grassy weeds, as confirmed by further evaluation with representative compounds with test dosages equivalent to an application rate of from 0.2 to 5.0 kg active ingredient, e.g. test dosages equivalent to a rate of 0.2, 1.0 and 5.0 kg active ingredient/ha, in dicotyledonous weeds such as *Amaranthus retroflexus*, *Capsella bursa-pastoris*, *Chenopodium alba*, *Stellaria media*, *Senecio vulgaris* and *Galium aparine*, and, especially, grassy weeds such as *Agropyron repens*, *Agrostis alba*, *Alopecurus myosuroides*, *Apera spica-venti*, *Avena fatua*, *Echinochloa crus-galli*, *Bromus tectorum*, *Sorghum halepense*, *Digitaria spp* and *Setaria spp*. Additional tests indicate a favourable soil persistence of the compounds of the invention.

The compounds of the invention are relatively less toxic towards crops than towards weeds. Selective herbicidal activity is i.a. observed in corn (maize), soybean, cotton, sugar beet, potato, alfalfa, sunflower, rape, peanuts or flax, depending i.a. on the compound involved and on the application rate. The compounds of the invention are therefore also indicated for use as selective herbicides in a crop locus.

The present invention therefore also provides a method of combatting weeds in a locus, preferably

in a crop locus, particularly in a crop locus as mentioned above, which comprises applying to the locus a herbicidally effective amount (a selective herbicidally effective amount where the locus is a crop locus) of a compound of the invention. A particularly preferred and advantageous embodiment of the invention is the pre-emergence (both crops and weeds) use of a compound of formula I in selectively combatting weeds in a crop locus.

For general herbicidal as well as for selective herbicidal use of compounds of the invention, the amount to be applied to attain the desired effect will vary depending on the particular crop if employed for selective use and other standard variables such as the compound employed, mode of application, conditions of treatment and the like. The appropriate application rates can be determined by routine procedures by those skilled in the art, or by comparing the activity of the compounds of the invention with standards for which the application rate is known, e.g. in greenhouse tests. However, in general, satisfactory results are usually obtained when the compound is applied at a rate in the range of from about 0.1 to 5 kg/ha, preferably from about 0.2 to 4 kg/ha, more preferably from 0.5 to 3.0 kg/ha, the application being repeated as necessary. When used in a crop locus, the application rate should preferably not exceed 3 kg/ha.

The compounds of formula I may be and preferably are employed as herbicidal compositions in association with herbicidally acceptable diluent(s). Suitable formulations contain 0.01% to 99% by weight of active ingredient, from 0 to 20% herbicidally acceptable surfactant and 1 to 99.99% solid or liquid diluent(s). Higher ratios of surfactant to active ingredient are sometimes desirable and are achieved by incorporation into the formulation or by tank mixing. Application forms of composition generally contain between 0.01 and 25% by weight of active ingredient. Lower or higher levels of active ingredient can, of course, be present depending on the intended use and the physical properties of the compound. Concentrate forms of composition intended to be diluted before use generally contain between 2 and 90%, preferably between 10 and 80% by weight of active ingredient.

Useful formulations of the compounds of the invention include dusts, granules, pellets, suspension concentrates, wettable powders, emulsifiable concentrates and the like. They are obtained by conventional manner, e.g. by mixing the compounds of the invention with the diluent(s). More specifically liquid compositions are obtained by mixing the ingredients, fine solid compositions by blending and, usually grinding, suspensions by wet milling and granules and pellets by impregnating or coating (preformed) granular carriers with the active ingredient or by agglomeration techniques.

Alternatively, the compounds of the invention may be used in microencapsulated form.

Herbicidally acceptable additives may be employed in the herbicidal compositions to improve the performance of the active ingredient and to reduce foaming, caking and corrosion.

Surfactant as used herein means a herbicidally

acceptable material which imparts emulsifiability, spreading, wetting, dispersibility or other surface-modifying properties. Examples of surfactants are sodium lignin sulphonate and lauryl sulphate.

- 5 Diluents as used herein mean a liquid or solid herbicidally acceptable material used to dilute a concentrated material to a usable or desirable strength. For dusts or granules it can be e.g. talc, kaolin or diatomaceous earth, for liquid concentrate
10 forms, for example a hydrocarbon such as xylene or an alcohol such as isopropanol, and for liquid application forms i.a. water or diesel oil.

- The compositions of this invention can also comprise other compounds having biological activity,
15 e.g. compounds having similar or complementary herbicidal activity or compounds having antidotal, fungicidal or insecticidal activity.

Specific Examples of herbicidal compositions will now be described.

20 EXAMPLE A: Wettable Powder

- 25 Parts of a compound of formula I, e.g. Compound No. 25 hereinafter given, are mixed and milled with 25 parts of synthetic fine silica, 2 parts of sodium lauryl sulphate, 3 parts of sodium ligninsulphonate and 45 parts of finely divided kaolin until the mean particle size is about 5 micron. The resulting
30 wettable powder is diluted with water before use to a spray liquor with the desired concentration.

EXAMPLE B: Emulsion Concentrate

- 30 20 Parts of a compound of formula I, e.g. Compound No. 25 hereinafter given, 40 parts of xylene, 30 parts of dimethyl formamide and 10 parts of emulsifier (e.g. ATLOX 4851 B a blend of Ca alkylarylsulphonate and a polyethoxylated triglyceride of Atlas Chemie GmbH) are thoroughly
35 mixed until a homogeneous solution is obtained. The resulting emulsion concentrate is diluted with water before use.

EXAMPLE C: Granules

- 40 5 Kg of a compound of formula I, e.g. Compound No. 25 hereinafter given, are dissolved in 25l methylene chloride. The solution is then added to 95 kg of granulated attapulgite (mesh size 24/48 mesh/inch) and thoroughly mixed. The solvent is then
45 evaporated off under reduced pressure.

The invention is further illustrated by the following Examples wherein temperatures are in °C, pressures are in Torr and Rf values are on silica gel, so far not otherwise indicated.

50 FINAL COMPOUNDS

Example 1 N - (2,4 - dimethyl - thien - 3 - yl) - N - (1,3 - dioxolane - 2 - ylmethyl) - chloroacetamide (process a)

- To the well stirred mixture of 1.7 g (0.0063 mol) of
55 N - (2,4 - dimethyl - thien - 3 - yl) - N - (1,3 - dioxolane - 2 - ylmethyl) - hydroxyacetamide, 58 mg of benzyldimethylammonium chloride, 7 ml of toluene and 7 ml of 30% NaOH are added dropwise without cooling 1.31 g (0.0069 mol) of p-toluene sulfonyl
60 chloride in 3.5 ml of toluene. After the exothermic reaction has subsided, stirring is continued for a further 20 hours at ambient temperature.

The toluene layer is then separated, washed with water and dried over anhydrous Na₂SO₄.

- 65 The crude residue left on rotevaporation of the

solvent is recrystallised from diethyl ether to give N - (2,4 - dimethyl - thien - 3 - yl) - N - (1,3 - dioxolane - 2 - ylmethyl) - tosyloxyacetamide ("tosylate") having a m.p. of 101-103°.

- 70 A stirred mixture of 2.16 g (0.0051 mol) of said tosylate, 1.84g (0.0066 mol) of tetrabutylammonium chloride, 8 ml of water and 16 ml of toluene is heated at 90° over a period of 6 hours. The toluene layer is then separated, dried (Na₂SO₄) and evaporated in
75 vacuo.

The residue is chromatographed on a silica gel column. Elution with hexane - ethylacetate 2:1 affords the title compound as analytically pure crystals, having a m.p. of 76-78°.

- 80 Example 2 N - (2,4 - dimethyl - thien - 3 - yl) - N - methoxyethyl - chloroacetamide (process a)

To a stirred solution of 2.0 g (0.008 mol) of N - (2,4 - dimethyl - thien - 3 - yl) - N - methoxyethyl -
85 hydroxyacetamid and 0.92 g (0.008 mol) of 4 - dimethylaminopyridine (=DMAP) in 80 ml of dry methylene chloride are added without cooling 0.93 g (0.008 mol) of mesyl chloride in 20 ml of dry CH₂Cl₂.

The resulting mixture containing DMAP-hydrochloride and the methanesulphonate of the starting compound is heated under reflux for 25 hours and then evaporated in vacuo.

The residual oil, after column chromatography on silica gel (elution with diethylether), solidified on
95 chilling at -20°, m.p. 54-55.

Example 3 : N - (chloroacetyl) - N - (2 - carbomethoxy - 4 - methyl - thien - 3 - yl) - alanine - ethylester (process b)

To a well stirred suspension of 1.5 g (0.05 mol)
100 sodium hydride (80% dispersion in mineral oil) in 250 ml of dry dimethoxyethane (=DME) are added portionwise, 9.9 g (0.04 mol) of solid N - (2 - carbomethoxy - 4 - methyl - thien - 3 - yl) - chloroacetamide.

105 After the addition is completed the resulting solution of the Na-salt is stirred an additional hour at 50°, then allowed to reach room temperature and treated with the solution of 7.25 g (0.04 mol) of ethyl - 2 - bromopropionate in 50 ml of dry DME.

110 After a reaction period of 4 hours at 50°, the mixture is filtered and evaporated in vacuo (50°/0.01 Torr.). The residual brown oil, after chromatography on silica gel (elution with diethyl ether-hexane 2:1), is subjected to ball tube distillation, giving the
115 analytically pure title compound, b.p. 135°/0.005 Torr.

Example 4 : N - (1H - pyrazol - 1 - ylmethyl) - N - (2,4 - dimethyl - thien - 3 - yl) - chloroacetamide (process b)

120 To a well stirred mixture of 19.35 g (0.095 mol) of N - (2,4 - dimethyl - thien - 3 - yl) - chloroacetamide, 4.15 g (0.01 mol) of benzyldimethylhexadecyl - ammonium chloride, 40 g (1 mol) of sodium hydroxide, 200 ml of methylene chloride and 40 ml of water are added 17 g (0.11 mol) of solid 1-chloromethyl
125 pyrazolehydrochloride at such a rate, that the temperature does not rise above 25°.

When the addition is completed, the reaction mixture is stirred an additional 2½ hours at ambient
130 temperature. Then 100 ml of water are added. The

organic layer is separated, washed with three 200 ml portions of water, dried over Na_2SO_4 and evaporated to dryness. The residue is chromatographed on a silica gel column. Elution with hexane-diethylether

5 1:1 affords the title compound as an analytically pure syrup which crystallized on chilling overnight at -20° , m.p. $88-89^\circ$ (recrystallized from diethyl ether).

Example 5 : N - (chloroacetyl) - N - (3,5 - dimethyl - isoxazol - 4 - yl) - alanineethylester
(process b)

10 9.4 g (0.05 mol) of N - (3,5 - dimethyl - isoxazol - 4 - yl) - chloroacetamide in 150 ml of dry acetonitrile ($=\text{CH}_3\text{CN}$) are added dropwise to a well stirred suspension of 1.8 (0.06 mol) of sodium hydride (80% dispersion in mineral oil) in 25 ml of dry CH_3CN .

15 After the exothermic reaction (34°) has subsided the solution of the sodium salt is allowed to reach room temperature and then treated with the solution of 9.05 g (0.05 mol) of ethyl 2 - bromopropionate in 25 ml of dry CH_3CN .

20 After the addition is completed the reaction mixture is heated at 50° for three hours and then evaporated to dryness. The residue is taken up with 100 ml of diethyl ether and filtered. The residual oil left on evaporating the filtrate is chromatographed on a silica gel column. Elution with diethylether-hexane 1:1 affords the title compound, having a m.p. of $49-50^\circ$.

Example 6 : N - [1 - (1H - pyrazol - 1 - yl) ethyl] - N - (2,4 - dimethyl - thien - 3 - yl) - chloroacetamide (process c)

30 To a stirred solution of 12.7 g (0.1 mol) of 2,4 - dimethyl - 3 - amino - thiophene in 100 ml of dry benzene, which contains 16 g of molecular sieves (3Å) and two drops of conc. H_2SO_4 , are added dropwise 8.5 ml (0.15 mols) of acetaldehyde, taking care that the temperature does not rise above 25° .

35 When all the acetaldehyde has been introduced, the reaction mixture is stirred 24 hours longer at ambient temperature and is then filtered. Removal of the solvent leaves the Schiff base as a light brown liquid.

40 To the stirred solution of 13 g (0.085 mols) of this material in 75 ml of dry toluene are added dropwise at -30° 9.7 g (0.086 mols) of chloroacetyl chloride.

45 When the addition is complete, the reaction solution is stirred for a further 30 minutes at -30° and then treated dropwise at the same temperature with 12 g (0.085 mols) of 1 - trimethylsilylpyrazole. After the addition, the solid CO_2 /acetone bath is removed, the mixture allowed to warm to ambient temperature and left to stir for a further period of 20 hours.

50 The reaction mixture is then filtered and evaporated in vacuo. The residual syrup is chromatographed on a silica gel column. Elution with hexane/diethyl ether 3:2 affords colourless crystals of the title compound which, upon recrystallization from diethyl ether have a melting point of $76-78^\circ$.

Example 7 : N - (4 - methoxy - 2 - methyl - thien - 3 - yl) - N - (2 - ethoxyethyl) - chloroacetamide (process d)

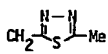
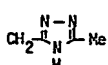
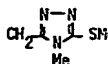





60 To a well stirred mixture of 6.05 g (0.03 mol) N - (2 - ethoxyethyl) - 4 - methoxy - 2 - methyl - thiophene - 3 - amine, 4.15 g (0.03 mol) of K_2CO_3 , 10 ml of water and 100 ml of CH_2Cl_2 is added dropwise without

cooling the solution of 3.4 g (0.03 mol) of chloroacetyl chloride in 10 ml of CH_2Cl_2 . After the exothermic reaction (27°) has subsided, stirring is continued for a further hour at ambient temperature.

70 The methylene chloride layer is then separated, washed twice with 100 ml of water, dried (Na_2SO_4) and evaporated in vacuo. The residual crude title compound is analytically pure; RF = 0.23 (hexane-diethyl ether 1:2). A small portion of this material was subjected to ball tube distillation: b.p. $168-170^\circ/0.05$ Torr.

75 The following compounds of formula I are obtained according to one or more of the procedures of the Examples 1 to 7, as indicated hereinbefore:
80 (Me is CH_3 and Et is C_2H_5)

Compd. No.	Aryl	Y	Characterization
1	4-Me-thien-3-yl	$\text{CH}_2\text{OC}_2\text{H}_5$	m.p. 25-26°
2	"	$\text{CH}_2\text{CH}_2\text{OCH}_3$	Rf=0.45(cyclohexane/ethylacetate 1:1)
3	2-Me-thien-3-yl	$\text{CH}_2\text{CH}_2\text{OCH}_3$	Rf=0.3(cyclohexane/ethylacetate 6:4)
4	2,4-diMe-thien-3-yl	CH_3	m.p. 45-46°
5	"	C_2H_5	m.p. 50-51°
6	"	tC_3H_7	
7	"	$\text{CH}_2=\text{CH}=\text{CH}_2$	
8	"	$\text{CH}_2\text{C}(\text{CH}_3)=\text{CH}_2$	m.p. 85-86°
9	"	$\text{CH}_2=\text{C}=\text{CH}$	m.p. 95-96°
10	"	$\text{C}(\text{CH}_3)_2-\text{C}=\text{CH}$	
11	"	$\text{CH}_2\text{C}_6\text{H}_5$	$n_D^{20}=1.5479$
12	"	CH_2CF_3	m.p. 60-61°
13	"	$\text{CH}_2-\text{C}(\text{Cl})=\text{CH}_2$	
14	"	$\text{CH}_2-\text{C}(\text{Br})=\text{CH}_2$	m.p. 45-46°
15	"	CH_2-COOH	m.p. 145-48°
16	"	CH_2COOMe	m.p. 48-49°
17	"	CH_2COOEt	$n_D^{20}=1.5345$
18	"	$\text{CH}_2\text{COOC}_3\text{H}_7$	b.p. 118-21°/0.005 Torr
19	"	$\text{CH}_2-\text{COOC}(\text{CH}_3)_2\text{C}=\text{CH}$	m.p. 77-79°
20	"	$\text{CH}(\text{CH}_3)\text{COOCH}_3$	$n_D^{20}=1.5342$
21	"	$\text{CH}(\text{CH}_3)\text{COOC}_3\text{H}_7$	$n_D^{20}=1.5192$
22	"	$\text{CH}_2\text{CON}(\text{CH}_3)_2$	m.p. 82-84°
23	"	$\text{CH}_2\text{CONHC}_3\text{H}_7$	m.p. 135-137°
24	"	$\text{CH}_2\text{CONHN}=\text{C}(\text{CH}_3)_2$	m.p. 141°
25	"	$\text{CH}_2-(1\text{-pyrazolyl})$	m.p. 88-89°
26	"	$\text{CH}(\text{CH}_3)(1\text{-pyrazolyl})$	m.p. 76-78°
27	"	$\text{CH}_2-(3,5\text{-diMe-pyrazolyl-1})$	m.p. 143-144°
Compd. No.	Aryl	Y	Characterization
28	2,4-diMe-thien-3-yl	$\text{CH}(\text{CH}_3)-\text{N} \begin{array}{c} \diagup \\ \diagdown \end{array}$	
29	"	$\text{CH}_2-\text{N} \begin{array}{c} \diagup \\ \diagdown \end{array}$	
30	"	$\text{CH}(\text{CH}_3)-\text{N} \begin{array}{c} \diagup \\ \diagdown \end{array}$	m.p. 132-133°
31	"	$\text{CH}_2-\text{N} \begin{array}{c} \diagup \\ \diagdown \end{array}$	
32	"	$\text{CH}(\text{Me})-\text{N} \begin{array}{c} \diagup \\ \diagdown \end{array}$	
33	"	$\text{CH}_2-\text{N} \begin{array}{c} \diagup \\ \diagdown \end{array}$	
34	"	$\text{CH}(\text{Me})-\text{N} \begin{array}{c} \diagup \\ \diagdown \end{array}$	
35	"	$\text{CH}(\text{Me})-\text{N} \begin{array}{c} \diagup \\ \diagdown \end{array}$	
36	"	$\text{CH}_2(2\text{-furyl})$	m.p. 78-79°
37	"	$\text{CH}_2(2\text{-thienyl})$	m.p. 57-59°
38	"	$\text{CH}_2-\text{N} \begin{array}{c} \diagup \\ \diagdown \end{array}$	m.p. 66-67°
39	"	3-Me-isoxazol-5-yl- CH_2	m.p. 77-78°
40	"	$\text{CH}_2-\text{N} \begin{array}{c} \diagup \\ \diagdown \end{array}$	m.p. 54-56°
41	"	$\text{CH}_2-\text{N} \begin{array}{c} \diagup \\ \diagdown \end{array}$	m.p. 71-72°

Compd. No.	Aryl	Y	Characterization
42	2,4-diMe-thien-3-yl		m.p. 110-15°
43	"		m.p. 119-25°
44	"		m.p. 120-22°
45	"		m.p. 68-70°
46	"	CH ₂ CH ₂ OH	m.p. 79-80
47	"	CH ₂ OEt	b.p. 115°/0.001 Torr
48	"	CH ₂ OC ₃ H ₇ n	n _D ²⁰ =1.5280
49	"	CH ₂ OC ₄ H ₉ n	b.p. 110-11°/0.001 Torr
50	"	CH(Me)OMe	m.p. 48-50°
51	"	CH(Et)OMe	m.p. 55-57°
52	"	CH ₂ CH ₂ OMe	m.p. 54-55°
53	"	CH ₂ CH ₂ OEt	b.p. 110°/0.01 Torr
54	"	CH ₂ CH ₂ OC ₃ H ₇ n	Rf=0.36(diethylether/hexane 1:1)
55	"	CH(Me)CH ₂ OMe	b.p. 148-150°/0.03 Torr
56	"	C(Me) ₂ CH ₂ OMe	
57	"		
58	"		m.p. 76-78°
Compd. No.	Aryl	Y	Characterization
59	2,4-diMe-thien-3-yl	CH(Me)OCH ₂ CH ₂ Cl	m.p. 62-64°
60	"	CH(OMe)CH ₂ OMe	Rf=0.3(diethylether/hexane 1:1)
61	"	CH ₂ OCH ₂ CH ₂ OMe	b.p. 117-18°/0.005 Torr
62	"	CH(Me)OCH ₂ CH ₂ CH ₃	m.p. 59-64°
63	"	CH ₂ OCH ₂ -N 	
64	"	CH(Me)OCH ₂ -N 	
65	"	CH(Me)O-N=CH-Me	
66	"	CH(Me)O-N=C(Me) ₂	
67	"	CH ₂ -CH=NOMe	Rf=0.31(hexane/ethyl-acetate 3:2)
68	"	CH ₂ -CH=NOEt	Rf=0.23(hexane/ethyl-acetate 4:1)
69	"	CH ₂ -C(Me)=NOMe	m.p. 76-78°
70	"	CH ₂ -C(Me)=NOEt	m.p. 57-59°
71	"	CH(Me)CH=NOH	m.p. 68-70°
72	"	CH(Me)CH=NOEt	Rf=0.4(hexane/ethyl-acetate 4:1)
73	"	CH(Me)-N(Me)-COMe	
74	2-Me-4-Et-thien-3-yl	CH(Me)CH ₂ OMe	b.p. 142-44°/0.2 Torr
75	"	CH ₂ OEt	m.p. 49-50°
76	"	pyrazolyl-1-CH ₂	m.p. 53-54°
77	2-Et-4-Me-thien-3-yl	CH ₂ OC ₂ H ₅	Rf=0.47(diethylether/hexane 7:3)
78	"	CH(Me)CH ₂ OMe	Rf=0.47(diethylether/hexane 7:3)
79	2,4-diEt-thien-3-yl	pyrazolyl-1-CH ₂	m.p. 63-65°
80	"	CH ₂ -OEt	n _D ²⁰ =1.5242
81	"	CH ₂ CH ₂ OMe	n _D ²³ =1.5328

Compd. No.	Aryl	Y	Characterization
82	2-(C ₃ H ₇) ₃ -4-Me-thien-3-yl	pyrazolyl-1-CH ₂	
83	4-OH-2-Me-thien-3-yl	CH(Me)CH ₂ OMe	
84	2-Me-4-MeO-thien-3-yl	pyrazolyl-1-CH ₂	m.p. 90-92°
85	"	CH ₂ -OEt	m.p. 24°
86	"	CH ₂ CH ₂ -OEt	b.p. 168-170°/0.05 Torr
87	2-Me-4-EtO-thien-3-yl	pyrazolyl-1-CH ₂	
88	2-Me-4-i-C ₃ H ₇ -thien-3-yl	"	
89	2-Me-4-nC ₄ H ₉ O-thien-3-yl	CH ₂ OEt	m.p. 44-46°
90	2-Me-4-OCH ₂ -CH=CH ₂ -thien-3-yl	pyrazolyl-1-CH ₂	
91	2-Me-4-OCH ₂ -C≡CH-thien-3-yl	"	
92	2-MeOCH ₂ -4-MeO-thien-3-yl	"	
93	2-MeS-4-Me-thien-3-yl	"	b.p. 180°/0.001 Torr
94	"	CH ₂ OEt	b.p. 135°/0.001 Torr
95	"	CH ₂ CH ₂ OMe	b.p. 148-150°/0.001 Torr
96	2-MeS(O)-4-Me-thien-3-yl	CH ₂ OEt	m.p. 100°
96a	2-MeSO ₂ -4-Me-thien-3-yl	CH ₂ OEt	
97	2-MeCO-4-Me-thien-3-yl	CH ₂ OEt	m.p. 37-38°
97a	2-MeCO-4-Et-thien-3-yl	pyrazolyl-1-CH ₂	
97b	2-MeCO-4,5-diMe-thien-3-yl	pyrazolyl-1-CH ₂	
98	2-MeC(=NOMe)-4-Me-thien-3-yl	CH ₂ OEt	syn : m.p. 89-91° anti : m.p. 75-76°

Compd. No.	Aryl	Y	Characterization
99	2-MeC(=NOMe)-4-Me-thien-3-yl	pyrazolyl-1-CH ₂	syn : m.p. 123°
100	2-MeC(OEt) ₂ -4-Me-thien-3-yl	CH ₂ OEt	m.p. 46-47°
101	2-Me-4-O-COMe-thien-3-yl	CH(Me)CH ₂ OMe	
102	2-COOMe-4-Me-thien-3-yl	CH ₂ -C≡CH	m.p. 119-21°
103	"	CH ₂ OEt	m.p. 20-22°
104	"	CH(Me)COOEt	b.p. 135°/0.005 Torr
105	2- <i>i</i> C ₃ H ₇ -4-COOMe-thien-3-yl	pyrazolyl-1-CH ₂	
106	2-Et-4,5-diMe-thien-3-yl	CH ₂ OEt	n _D ²⁰ = 1.5273
107	"	CH ₂ COOC ₃ H ₇	n _D ²⁰ = 1.5112
108	"	pyrazolyl-1-CH ₂	n _D ²⁰ = 1.5509
109	2,4-diMe-5-Cl-thien-3-yl	CH ₂ OEt	n _D ²⁰ = 1.5412
110	"	CH ₂ OCH ₂ CH ₂ OCH ₃	n _D ²¹ = 1.5321
110a	"	tetrahydrofuryl-2-CH ₂	Rf=0.35(cyclohexane/ethylacetate 7:3)
110b	"	<i>i</i> C ₃ H ₇	Rf=0.38(cyclohexane/ethylacetate 6:4)
111	"	pyrazolyl-1-CH ₂	m.p. 68-73°
112	2,5-diBr-4-Me-thien-3-yl	CH ₂ OEt	m.p. 75-77°
113	2-Me-4-MeO-5-Br-thien-3-yl	pyrazolyl-1-CH ₂	m.p. 98-99°
114	2,4-diMe-5-COOMe-thien-3-yl	CH ₂ OEt	b.p. 140°/0.005 Torr
115	2,4-diMe-furan-3-yl	CH ₂ OEt	Rf=0.5(diethylether)

Compd. No.	Aryl	Y	Characterization
116	2,4-diMe-furan-3-yl	CH ₂ CH ₂ OMe	
117	"	CH ₂ -CH=NOMe	
118	"	CH ₂ -C(Me)=NOMe	
119	"	pyrazolyl-1-CH ₂	
120	1,2,4-triMe-pyrrol-3-yl	CH ₂ -OEt	
121	"	CH ₂ CH ₂ OMe	
122	"	pyrazolyl-1-CH ₂	
123	"	CH ₂ -C(Me)=CH ₂	
124	2-COOEt-N,3,5-tri-Mepyrrol-4-yl	pyrazolyl-1-CH ₂	n _D ²⁴ =1.5422
125	3,5-diMe-isoxazol-4-yl	CH(Me)COOEt	m.p. 49-50°
126	"	CH ₂ CH ₂ OMe	
127	"	CH ₂ OEt	m.p. 45-46°
128	"	pyrazolyl-1-CH ₂	
129	3,5-diEt-isoxazol-4-yl	CH ₂ -C(Me)=CH ₂	
130	"	CH ₂ -C≡CH	b.p. 118°/0.001 Torr
131	"	CH ₂ -OEt	{ b.p. 107-108°/0.001 Torr n _D ²⁰ =1.4908
132	3,5-diMe-isothiazol-4-yl	CH ₂ OEt	m.p. 43-45°
133	"	CH ₂ CH ₂ OMe	
134	"	pyrazolyl-1-CH ₂	m.p. 109-14°
135	"	CH ₂ C≡CH	m.p. 109-12°
136	3-Me-5-EtO-pyrazol-4-yl	CH ₂ CH ₂ OMe	m.p. 111-113°
137	1,3,5-triMe-pyrazol-4-yl	CH ₂ OEt	b.p. 130°/0.001 Torr
Compd. No.	Aryl	Y	Characterization
138	1,3,5-triMe-pyrazol-4-yl	CH ₂ CH ₂ OMe	m.p. 66-67°
139	"	CH ₂ -OC ₃ H ₇ n	b.p. 135°/0.001 Torr
140	"	Et	m.p. 80-82°
141	"	CH ₂ -C≡CH	m.p. 115-117°
142	"	pyrazolyl-1-CH ₂	m.p. 96-97°
143	1-Me-3,5-diEt-pyrazol-4-yl	CH ₂ -OC ₃ H ₇ n	n _D ²⁰ =1.5008
144	"	CH ₂ CH ₂ OMe	b.p. 130°/0.001 Torr
145	1,3-diMe-5-EtO-pyrazol-4-yl	CH ₂ OEt	m.p. 54-56°

INTERMEDIATES

Example 8 : Methyl 3,5 - dimethyl - 4 - amino - thiophene - 2 - carboxylate

A solution of 45.2 g (0.21 mol) methyl 3,5 - dimethyl - 4 - nitrothiophene - 2 - carboxylate in 1000 ml methanol is hydrogenated for two hours at 10 bar in the presence of 4.5 g palladium (10% on carbon).

When the reduction is completed the mixture is filtered, the catalyst washed with methanol and the filtrate evaporated to dryness.

The crystalline residue is treated with diethyl ether, yielding the analytically pure title compound m.p. 88-89°

Example 9 : 2,4-Dimethyl-3-aminothiophene

To 890 ml (3 mol) of sodium bis(2 - methoxyethoxy) aluminium hydride (70% solution in toluene)

and 600 ml of dry toluene is added, dropwise with vigorous stirring, a solution of 100 g (0.58 mol) of methyl 3 - amino - 4 - methyl - thiophene - 2 - carboxylate in 700 ml of dry toluene at such a rate that the temperature does not rise above 55°.

After the addition is completed stirring is continued for a further 30 minutes and the reaction mixture then cautiously added in small portions at 0° to 1200 ml of 20% potassium hydroxide solution.

The toluene layer is separated, dried over MgSO₄ and evaporated in vacuo. The residual brown liquid is distilled under diminished pressure, affording the analytically pure title compound, b.p. 49-52°/0.01 Torr.

Example 10 : 4 - Methoxy - 2 - methyl - 3 - amino - thiophene

The well stirred mixture of 55.4 g (0.2 mol) of benzyl N - (4 - methoxy - 2 - methyl - thien - 3 - yl) carbamate, 40 g (0.7 mol) of KOH, 600 ml of ethanol and 120 ml of water is heated under reflux for two

- 5 hours. The resulting solution is then evaporated in vacuo and the residue diluted with 500 ml of water. The separating light yellow oil is taken up in 400 ml of diethyl ether and the aqueous phase extracted once with 400 ml of ether.
- 10 The combined ethereal solutions are dried (Na_2SO_4) and filtered. A slow current of dry hydrogen chloride is then passed through the filtrate for a period of 30 minutes, taking care - by
- 15 intermittent colling in an ice bath - that the temperature does not rise above 10° .

The precipitated hydrochloride of the title compound has a m.p. of 230° (the free base a m.p. of $61-63^\circ$).

Example 11 : 2 - Methylthio - 4 - methyl - 3 - aminothiophene

- 20 6 g (0.22 gram atoms) freshly prepared aluminium amalgam are added over a period of about two minutes to 18.9 g (0.1 mol) of 2 - methylthio - 4 - methyl - 3 - nitrothiophene dissolved in 200 ml of
- 25 moist diethylether. The vigorous reaction which sets in after 5-10 minutes is kept under control by cooling the flask in ice.

- After 45 minutes, when the exothermic reaction has subsided and all the aluminium has reacted,
- 30 another six grams of amalgamated aluminium are added.

- The reaction mixture is then gently refluxed for 2 hours; during this period the formation of the thiophene amine is complete. The organic phase is
- 35 decanted and the remainder washed with two 50 ml portions of diethylether. The combined ethereal solutions are dried (MgSO_4) and evaporated in vacuo. The residual crude title compound is distilled under reduced pressure, having a boiling point of
- 40 $79-81^\circ/0.5$ Torr.

Example 12

In the following table are listed compounds of formula IX that are obtained according to one or more of the procedures of Examples 8 to 11.

- 45
- | | |
|----------|--|
| Ex. 12.1 | 1,3 - di - CH_3 - 5 - OC_2H_5 - 4 - NH_2 pyrazole (Rf = 0.28 tetrahydrofuran : hexane 9:1) |
| Ex. 12.2 | 3,5 - di - CH_3 - 4 - NH_2 - isothiazole (m.p. $82-84^\circ$) |
| Ex. 12.3 | 4 - C_2H_5 - 2 - CH_3 - 3 - aminothiophene, Rf = 0.25 (diethylether: hexane 2:1) |
| Ex. 12.4 | 4 - n - butoxy - 2 - methyl - 3 - amino - thiophene, m.p. 25° ; (Rf = 0.25 (diethylether/hexane 2:1) |
| Ex. 12.5 | 5 - $\text{C}_2\text{H}_5\text{O}$ - 3 - CH_3 - 4 - NH_2 - pyrazole m.p. $81-83^\circ$ |
| Ex. 12.6 | 3,5 - di - C_2H_5 - 1 - CH_3 - 4 - NH_2 - pyrazole b.p. $68-70^\circ/0.001$ Torr |
| Ex. 12.7 | 3,5 - di - C_2H_5 - 4 - NH_2 - isoxazole Rf=0.39 (diethyl ether) |
| Ex. 12.8 | (3 - NH_2 - 4 - Me - thien - 2 - yl)ethanone, m.p. $80-81^\circ$ |
| Ex. 12.9 | (3 - NH_2 - 4 - Et - thien - 2 - yl)ethanone, m.p. 68° |
- 65

Ex. 12.10 (3 - NH_2 - 4,5 - diMe - thien - 2 - yl)ethanone, m.p. $120-23^\circ$

Ex. 12.11 2 - Et - 4 - Me - 3 - NH_2 - thiophene, b.p. $74-76^\circ/0.01$ Torr

70 Ex. 12.12 2,4 - diEt - 3 - NH_2 - thiophene, Rf = 0.24 (CH_2Cl_2) $n_D^{20} = 1.5511$

Ex. 12.13 2 - Et - 4,5 - diMe - 3 - NH_2 - thiophene, $n_D^{20} = 1.5581$

75 *Example 13 : N - (2,4 - dimethyl - thien - 3 - yl) - acetoxyacetamide*

To 12.7 g (0.1 mol) of 2,4 - dimethyl - 3 - aminothiophene, 13.8 g (0.1 mol) of K_2CO_3 , 20 ml of water and 150 ml of methylene chloride are added dropwise at ambient temperature 15 g (0.11 mol) of acetoxyacetyl chloride.

- 80 After the exothermic reaction has subsided, stirring is continued for a further hour. Then the organic layer is evaporated, washed with water, dried
- 85 (MgSO_4) and evaporated in vacuo. The solid residue is treated with hexane affording the analytically pure title compound, m.p. $110-112^\circ$.

Example 14 : N - (2,4 - dimethyl - thien - 3 - yl) - hydroxyacetamide

- 90 Through a stirred solution of 15.9 g (0.07 mol) of N - (2,4 - dimethyl - thien - 3 - yl) - acetoxyacetamide in 300 ml of methanol is bubbled a gentle stream of ammonia gas for 30 minutes.

- After the exothermic (40°) reaction (moderated by
- 95 intermittent cooling in cold water) has subsided, stirring is continued for additional 30 minutes and the reaction solution then evaporated to dryness. The resulting solid is recrystallized from ethylacetate, giving the analytically pure title compound as
- 100 colourless crystals, m.p. $85-87^\circ$.

Example 15 : N - (2 - ethoxyethyl) - 4 - methoxy - 2 - methyl - thiophene - 3 - amine

- To 70 ml (0.245 mol) of sodium bis (2 - methoxyethoxy) aluminium hydride (70% solution in toluene)
- 105 and 30 ml of dry toluene is added dropwise with stirring a solution of 9.4 g (0.041 mol) of N - (4 - methoxy - 2 - methyl - thien - 3 - yl) - ethoxyacetamide in 125 ml of dry toluene. After the exothermic reaction has subsided, stirring is continued for a
- 110 further 90 minutes and the reaction solution then cautiously added in small portions at -10° to 100 ml of 20% KOH solution.

- The toluene layer is separated and the aqueous phase extracted twice with 100 ml of toluene. The
- 115 combined toluene solutions are dried (Na_2SO_4) and evaporated in vacuo. The residual light brown liquid is analytically pure, Rf = 0.32 (hexane - diethyl ether 1:2).

Example 16 : N - [(Methoxymethyl)ethyl] - 2,4 - dimethyl - thiophene - 3 - amine

- The solution of 6.4 g (0.05 mol) of 2,4 - dimethyl - 3 - amino - thiophene and 5.3 g (0.06 mol) of methoxyacetone in 100 ml of dry toluene is boiled under reflux until the theoretical amount of water has
- 125 separated in the water trap ($2\frac{1}{2}$ hours). The toluene solution is then allowed to cool to ambient temperature and evaporated in vacuo. The residual crude Schiff base is sufficiently pure for the next step.

- 9.2 g (0.048 mol) of this material are dissolved in
- 130 100 ml of dry toluene and added without cooling to a

well stirred solution of 28 ml (0.098 mol) of sodium bis (2 - methoxyethoxy) aluminium hydride (70% solution in toluene) in 15 ml of dry toluene.

- After the exothermic reaction (44°) has subsided, stirring is continued for a further 90 minutes and the brown reaction solution then cautiously added in small portions at -10° to 40 ml of 20% KOH solution.

- The toluene layer is then separated, dried (Na₂SO₄) and evaporated in vacuo. The residual oil, after column chromatography on silica gel (elution with hexane - diethyl ether 3:1) is distilled under diminished pressure (ball tube), having a b.p. of 94-95°/0.01 Torr.

- Example 17 : N - (5 - Ethyl - 1,3,4 - oxdiazol - 2 - ylmethyl) - 2,4 - dimethyl - thiophene - 3 - amine*

- The mixture of 12.7 g (0.1 mol) of 2,4 - dimethyl - 3 - amino - thiophene, 14.6 g (0.1 mol) of 2 - chloromethyl - 5 - ethyl - 1,3,4 - oxdiazole and 13.8 g (0.1 mol) of K₂CO₃ in 150 ml of dry dimethylformamide (=DMF) is stirred for 20 hours at 80°.

- The reaction mixture is then cooled to ambient temperature poured into 400 ml of water and the water-DMF solution extracted with three 150 ml portions of diethyl ether. The combined ethereal extracts are dried over anhydrous Na₂SO₄ and evaporated in vacuo. The residual brown oil is chromatographed on a silica gel column, elution with hexane/diethyl ether 1:1 affords the title compound as an analytically pure orange liquid, Rf=0.2 (ether-hexane 1:1).

- Example 18 : N - (2,4 - dimethyl - thien - 3 - yl)glycine hydrazide*

- 82 g (0.412 mol) N - (2,4 - dimethyl - thien - 3 - yl) glycine - methylester and 42 g (1.3 mol) hydrazine hydrate in 350 ml alcohol are stirred for 24 hours at room temperature. The mixture is evaporated in vacuo. After adding 150 ml water the residue is treated with 4 x 500 ml portions of ether. The ether phase is dried over Na₂SO₄ and evaporated to dryness giving the analytically pure title compound, m.p. 68-70°C

- Example 19 : N - (2,4 - dimethyl - thien - 3 - yl) - N - (4 - methyl - 5 - mercapto - 1,2,4 - triazol - 3 - yl - methyl) - amine*

- A mixture of 15 g (0.075 mol) N - (2,4 - dimethyl - thien - 3 - yl) glycine hydrazide and 5.5g (0.075 mol) methylisothiocyanate in 100 ml ethanol is refluxed for 4 hours. The reaction mixture is cooled to 6°C and the precipitate formed separated by filtration giving the pure compound, m.p. 184-87°C.

- Example 20 : N - (2,4 - dimethyl - thien - 3 - yl) - N - (4 - methyl - 5 - methylmercapto - 1,2,4 - triazol - 3 - yl - methyl) - amine*

- To a mixture of 9.5 g (0.037 mol) N - (2,4 - dimethyl - thien - 3 - yl) - N - (4 - methyl - 5 - mercapto - 1,2,4 - triazol - 3 - yl - methyl) - amine, 1.1g triethyl - benzyl ammonium chloride in 100 ml toluene and 30 ml aqueous 50% sodium hydroxide solution are added 4.7g (0.037 mol) dimethyl-sulfate. The reaction mixture is stirred at room temperature for 5 hours, the organic layer is separated, washed with three 100 ml portions of water, dried over Na₂SO₄ and evaporated to dryness. The residue is recrystallized twice from ether giving the pure title compound,

m.p. 92-96°C.

Example 21 : N - (2,4 - dimethyl - thien - 3 - yl) - N - (5 - methyl - 1,3,4 - thiadiazol - 2 - yl - methyl) - amine

- To 10g (0.05 mol) N - (2,4 - dimethyl - thien - 3 - yl) - glycine hydrazide in 175 ml methylene chloride is added dropwise 5.5g (0.05 mol) acetic acid anhydride. The mixture is stirred for 1 hour at 20°C and evaporated to dryness. The oily residue is dissolved in 150 ml pyridine and 11.1g (0.05 mol) phosphorous pentasulfide is added in small portions. During the course of the addition the temperature reached 48°C. The mixture is refluxed for two hours and stirred at room temperature for another 16 hours. After evaporation to dryness the residue is dissolved in methylene chloride and the solution treated successively with ice water, diluted cold sodium hydroxide solution and water, dried over Na₂SO₄ and evaporated to dryness. Chromatography on a silica gel column afforded the pure compound as an oil, Rf = 0.23 (ethylacetate/hexane 6:4).

Example 22 : N - (2,4 - dimethyl - thien - 3 - yl) - N - (5 - methyl - 1,2,4 - triazol - 3 - yl - methyl) amine



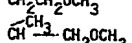
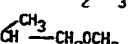



- To a stirred solution of 2.2g (0.055 mol) NaOH in 125 ml methanol is added 5 g (0.05 mol) acetamidine hydrochloride. After 15 minutes 10 g (0.05 mol) N - (2,4 - dimethyl - thien - 3 - yl) - glycine hydrazide is added and the mixture refluxed for 18 hours. After evaporation to dryness the residue is treated with methylene chloride and freed from insoluble by-products by filtration. Evaporation of the methylene chloride gives the title compound, m.p. 85-87°C.

Example 23 : N - (2,4 - dimethyl - thien - 3 - yl) glycine - isopropylidene - hydrazide

- 7g (0.035 mol) N - (2,4 - dimethyl - thien - 3 - yl) - glycine hydrazide in 50 ml acetone is stirred at room temperature for 2 hours. The pure title compound which crystallized was obtained by filtration, m.p. 126-28°C.

Example 24

The following compounds of formula VII are obtained according to one or more of the preceding Examples 13 to 23:

Ex.	Ar	Y	Characteriza- tion
24.1	2,4-di-CH ₃ -thien-3-yl	CH ₂ CH ₂ -OCH ₃	Rf=0.32(diethyl ether-hexane 3:1)
24.2	"	C ₂ H ₅	Rf=0.23(diethyl ether-hexane 1:1)
24.3	"	CH ₂ 	Rf=0.38(diethyl ether-hexane 1:1)
24.4	"	CH ₂ 	Rf=0.39(diethyl ether-hexane 2:1)
24.5	2-CH ₃ S-4-CH ₃ -thien-3-yl	CH ₂ CH ₂ -OCH ₃	b.p. 86°/0.01 Torr
24.6	1,3,5-tri-CH ₃ -pyrazol-4-yl	"	b.p. 92-94°/ 0.01 Torr
24.7	1-CH ₃ -3,5-di-C ₂ H ₅ - pyrazol-4-yl	"	Rf=0.38(acetone- hexane 4:6)
24.8	1,3,5-tri-CH ₃ -pyrazol-4-yl	C ₂ H ₅	Rf=0.2(ethyl- acetate-CH ₃ OH 2:1)
24.9	3-CH ₃ -5-OC ₂ H ₅ -pyrazol-4-yl	CH ₂ CH ₂ OCH ₃	Rf=0.25(ethyl- acetate-CH ₃ OH 9:1)
24.10	2,4-di-CH ₃ -thien-3-yl	CH ₂ CH ₂ OC ₂ H ₅	Rf=0.45(diethyl ether)
24.11	"	CH ₂ CH ₂ OC ₃ H ₇ -n	Rf=0.57(diethyl ether)
24.12	"	CH ₂ CF ₃	n _D ²⁰ =1.4786
24.13	2,4-di-C ₂ H ₅ -thien-3-yl	CH ₂ CH ₂ OCH ₃	n _D ²⁰ =1.5238
24.14	2-CH ₃ -4-C ₂ H ₅ -thien-3-yl		b.p. 98-100°/ 0.09 Torr
24.15	2-C ₂ H ₅ -4-CH ₃ -thien-3-yl		Rf=0.52 (ether-hexane 7:3)
24.16	2,4-di-CH ₃ -thien-3-yl	CH ₂ -C(CH ₃)=CH ₂	Rf=0.55(diethylether- hexane 1:1)
24.17	"	CH ₂ 	
24.18	"	CH ₂ 	
24.19	"	CH ₂ 	Rf=0.15(CH ₂ Cl ₂)
24.20	"	CH ₂ CON(CH ₃) ₂	n _D ²⁰ =1.5579
24.21	"	CH ₂ COOCH ₃	b.p. 82-87°/0.005 Torr

Example 25 : N - (2 - carbomethoxy - 3,5 - dimethyl - thien - 4 - yl) - chloroacetamide

To 12.25 g (0.066 mol) of methyl 3,5 - dimethyl - 4 - aminothiophene - 2 - carboxylate, 9.1 g (0.066 mol) of K₂CO₃, 25 ml of water and 150 ml of methylene chloride are added without cooling 7.5 g (0.066 mol) of chloroacetyl chloride.

After the exothermic reaction (36°) has subsided stirring is continued for a further hour. Then the organic layer is separated, washed with water and dried over anhydrous Na₂SO₄. After removal of the solvent the desired product is obtained as colourless crystals, m.p. 157-58°.

Example 26 : N - (2 - chloro - 3,5 - dimethyl - thien - 4 - yl) - chloroacetamide

To the stirred solution of 5.0g (0.0245 mol) of N - (2,4 - dimethyl - thien - 3 - yl) - chloroacetamide in 50 ml of dry CH₂Cl₂ are added dropwise at 0° 3.3g (0.0245 mol) of sulfonyl chloride.

When the addition is complete, the reaction mixture is allowed to warm to ambient temperature and

stirred for a further period of 20 hours. Then the solvent is removed in vacuo and the crystalline residue triturated with hexane, yielding the analytically pure title compound, m.p. 166°, decomp.

Example 27 : N - [2 - (1,1 - diethoxy) ethyl - 4 - methyl - thien - 3 - yl] - chloroacetamide

15 g (0.065 mol) of N - (2 - acetyl - 4 - methyl - thien - 3 - yl) - chloroacetamide are dissolved in the warm (50°) mixture of 60 g of dry ethyl alcohol, 60 g of triethyl orthoformate and 7 drops of concentrated HCl.

Upon standing for 30 hours at room temperature the reaction mixture is evaporated in vacuo (50°/0.01 Torr) and the residual brown syrup chromatographed on a silica gel column. Elution with hexane-diethylether 2:1 affords the title compound as colourless crystals, having a m.p. of 57-58°.

Example 28 : N - [2 - (1 - methoxyimino) ethyl - 4 - methyl - thien - 3 - yl] - chloroacetamide

The solution of 1.5 g (0.0065 mol) of N - (2 - acetyl -

4 - methyl - thien - 3 - yl) - chloroacetamide and 6.6 g (0.14 mol) of methoxyamine in 50 ml of dry toluene, which contains 5 g of molecular sieves (3Å) is refluxed for 7 hours.

- 5 The yellow reaction solution is then filtered and evaporated in vacuo. The resulting crude syn/anti isomer mixture is separated by chromatography with hexane-diethylether as the mobile phase.

First are eluted 0.6g of the pure isomer (syn isomer) with the m.p. of 101-102° (Rf=0.33, hexane-ether 1:1). Continued elution of the silica gel column affords the other analytically pure anti isomer (0.4 g) having a melting point of 87-89° (Rf=0.21).

15 Example 29

The following compounds of formula III are obtained according to one or more of the Examples 25 to 28:

Ex.	Ar	Characterization
29.1	2-COOCH ₃ -4-CH ₃ -thien-3-yl	m.p. 118-119°
29.2	2-CH ₃ -4-CH ₃ -thien-3-yl	m.p. 105-106°
29.3	2,4-di-CH ₃ -thien-3-yl	m.p. 128-129°
29.4	3,5-di-CH ₃ -isoxazol-4-yl	m.p. 96-98°
29.5	3,5-di-C ₂ H ₅ -isoxazol-4-yl	m.p. 67-69°
29.6	4-CH ₃ -thien-3-yl	m.p. 93-96°
29.7	2-CH ₃ -4-C ₂ H ₅ -thien-3-yl	m.p. 114°
29.8	2-CH ₃ -4-OC ₂ H ₅ -thien-3-yl	m.p. 144-45°
29.9	2,4-di-CH ₃ -furan-3-yl	m.p. 95-97°
29.10	3,5-di-CH ₃ -isothiazol-4-yl	m.p. 104-06°
29.11	3-CH ₃ -5-OC ₂ H ₅ -pyrazol-4-yl	m.p. 125-27°
29.12	2-COOC ₂ H ₅ -N,3,5-tri-CH ₃ -pyrrol-4-yl	m.p. 163-65°
29.13	2-C ₂ H ₅ -4-CH ₃ -thien-3-yl	m.p. 105°
29.14	2,4-di-C ₂ H ₅ -thien-3-yl	m.p. 145°
29.15	2-COCH ₃ -4-CH ₃ -thien-3-yl	m.p. 110°
29.16	2-CH ₃ -4-OC ₄ H ₉ -thien-3-yl	m.p. 129-30°
29.17	2-C ₂ H ₅ -4,5-di-CH ₃ -thien-3-yl	m.p. 147-48°

20 Example 30 : N - (2,4 - dimethylthien - 3 - yl) - N - methoxyethyl - hydroxyacetamide

18.4 g (0.1 mol) of N - (2,4 - dimethylthien - 3 - yl) - hydroxyacetamide in 50 ml of dry dimethylformamide (=DMF) are added dropwise to a well stirred suspension of 3.0 g (0.1 mol) of sodium hydroxide (80% dispersion in mineral oil) in 50 ml of dry DMF.

25 After the exothermic reaction (50°) has subsided, the solution of the Na-salt is allowed to reach room temperature and then treated with the solution of 10.4 g (0.11 mol) of 2 - chloroethylmethylether in 20 ml of dry DMF. When the addition is completed, the resulting mixture is heated at 100° for 4 hours and then evaporated to dryness (40°/0.1 Torr).

30 The residue is taken up with 200 ml of diethyl ether, washed with 250 ml of water, dried (MgSO₄) and filtered. The residual oil left on evaporating the filtrate is chromatographed on a silica gel column. Elution with ether affords the desired product as an analytically pure viscous liquid (Rf=0.25/ether).

Example 31

40 Following the procedure of Example 30, the following compounds of formula II, in which Ar is 2,4 - dimethylthien - 3 - yl, are obtained.

Ex.	Y	Characterization
31.1	CH ₂ CH=NOCH ₃	Rf=0.21 (hexane-ethyl-acetate 3:2)
31.2	CH ₂ CH=NOC ₂ H ₅	Rf=0.25 (hexane-ethyl-acetate 3:2)
31.3	CH ₂ C=NOCH ₃	m.p. 69-71°
31.4	CH ₂ -C(=NOCH ₃)-CH ₃	Rf=0.27 (hexane-ethyl-acetate 3:2)
31.5	CHCH=NOCH ₃	Rf=0.31 (hexane-ethyl-acetate 3:2)
31.6	CHCH=NOC ₂ H ₅	Rf=0.3 (hexane-ethyl-acetate 3:2)
31.7	CH ₂ -C(=O)-CH ₃	Rf=0.12 (hexane-ethyl-acetate 3:2)

Example 32 : N - (4 - Methoxy - 2 - methyl - thien - 3 - yl) ethoxyacetamide

45 To 9.8 g (0.068 mol) 4 - methoxy - 2 - methyl - 3 - amino - thiophene, 9.45 g (0.068 mol) K₂CO₃, 35 ml of water and 100 ml of CH₂Cl₂ is added without cooling the solution of 8.4 g (0.069 mol) of ethoxyacetyl chloride in 20 ml of CH₂Cl₂.

50 After the exothermic reaction (32°) has subsided, stirring is continued for a further hour. Then the organic layer is separated, washed twice with 150 ml of water and dried over anhydrous Na₂SO₄.

55 The residue left on rotevaporation of the solvent is subjected to column chromatography on silica gel. Elution with hexane - diethylether 1:1 affords the analytically pure title compound having a m.p. of 40-41°.

Example 33

60 Analogous to the procedure of Example 34, the following amides of formula ArNHCOY₃ are obtained.

Ex.	Ar	Y ₃	Characterization
33.1	2,4-di-CH ₃ -thien-3-yl	-CH ₂ -OCH ₃	m.p. 62-63°
33.2	"	-CH ₃	m.p. 168-69°
33.3	"	-C(=O)-CH ₃	m.p. 140-42°
33.4	"	-C(=O)-C ₂ H ₅	m.p. 164-65°
33.5	1,3,5-tri-CH ₃ -pyrazol-4-yl	-CH ₂ OCH ₃	m.p. 85-87°
33.6	1-CH ₃ -3,5-di-C ₂ H ₅ -pyrazol-4-yl	"	b.p. 120°/0.001 Torr
33.7	3-CH ₃ -5-OC ₂ H ₅ -pyrazol-4-yl	-CH ₂ OCH ₃	m.p. 123-30°
33.8	2,4-di-CH ₃ -thien-3-yl	-CH ₂ OC ₂ H ₅	m.p. 37-38°
33.9	"	-CH ₂ OC ₃ H ₇ -n	m.p. 43-45°
33.10	"	-CF ₃	m.p. 94-97°
33.11	2,4-di-C ₂ H ₅ -thien-3-yl	-CH ₂ OCH ₃	m.p. 48-49°

Example 34 : Benzyl (N - 4 - methoxy - 2 - methyl - thien - 3 - yl) carbamate

65 63.6 g (0.23 mol) of diphenylphosphoryl azide and 24.4 g (0.23 mol) of triethylamine are added all at once to the stirred suspension of 36.5 g (0.21 mol) of 4 - methoxy - 2 - methyl - thiophene - 3 - carboxylic acid in 300 ml of dry benzene.

70 The resulting mixture is refluxed for one hour and then treated with 25 g (0.23 mol) of benzyl alcohol.

After a reaction period of 3½ hours at 78° the reaction mixture is cooled to ambient temperature,

diluted with diethyl ether (350 ml) and washed successively with 250 ml of 5% HCl, 250 ml of saturated NaHCO₃ solution and 200 ml of saturated NaCl solution. The crystalline residue left on reevaporation of the dried (Na₂SO₄) solvent is triturated with pentane, yielding the analytically pure title compound, m.p. 107-108°.

Example 35

Following the procedure of Example 34, employing the appropriate carboxylic acid, the following carbamate is obtained.

Ex. 35.1 Benzyl (N - 2,4 - di - CH₃ - furan - 3 - yl) carbamate, m.p. 103-5°.

Ex. 35.2 Benzyl (N - 4 - n - butoxy - 2 - methyl - thien - 3 - yl) carbamate, m.p. 83°.

Example 36 : 4 - Methoxy - 2 - methyl - thiophene - 3 - carboxylic acid

The title compound is obtained by saponification (KOH) of the corresponding ethyl ester; m.p. 127°.

Example 37

Analogous to the procedure of Example 36 is obtained 4 - n - butoxy - 2 - methyl - thiophene - 3 - carboxylic acid, m.p. 66-68°.

Example 38 : Ethyl 4 - methoxy - 2 - methyl - thiophene - 3 - carboxylate

20 ml of 50% sodium hydroxide solution are added dropwise to a mixture of 18.6 g (0.1 mol) of ethyl 4 - hydroxy - 2 - methyl - thiophene - 3 - carboxylate, 10 ml (0.105 mol) of dimethyl sulphate and 2.3 g (0.01 mol) of benzyltriethylammonium chloride in 100 ml of CH₂Cl₂, at ambient temperature.

After about 15 minutes of reflux, the reaction mixture is worked up to give the title compound, Rf = 0.36 (diethyl ether:hexane 2:1).

Example 39 : Methyl 3,5 - dimethyl - 4 - nitrothiophene - 2 - carboxylate

To a well stirred, chilled (0°) solution of 51.4 g (0.3 mol) of methyl 3,5 - dimethylthiophene - 2 - carboxylate in 200 ml of glacial acetic acid are added dropwise over a period of 40 minutes, a mixture of 30 ml of fuming nitric acid (specific gravity = 1.5) and 120 ml of acetic anhydride. After the addition is completed the resulting brown solution is stirred a further two hours at 5° and then poured into 3000 ml of ice water.

The aqueous phase is extracted twice with 500 ml of diethyl ether. The ethereal extracts are washed with water and 3% sodium bicarbonate solution, dried (Na₂SO₄) and evaporated in vacuo.

The residue is subjected to column chromatography on silica gel. Elution with hexane - diethylether (10:1) afforded the title compound having a melting point of 87-88°.

Example 40

Analogous to the process of Example 39 but employing 1,3 - dimethyl - 5 - ethoxypyrazole as starting material is obtained 1,3 - dimethyl - 5 - ethoxy - 4 - nitropyrazole; Rf = 0.37 with diethylether on silica gel.

Herbicidal Tests

Example 41: Weed control - Pre-emergence treatment

Seed pots (7 cm diameter) are filled with a mixture of peat culture substrate and sand. The exposed surface of the peat culture substrate and sand

mixture is sprayed with a test liquid of a test compound (e.g. formulated in accordance with Example B) and seeds of *Lepidium sativum*, *Agrostis alba*, *Avena sativa* and *Lolium perenne* are sown in each pot, whereby the *Avena sativa* and *Lolium perenne* seeds are, after sowing covered with a thin layer (0.5 cm) of peat culture substrate/sand mixture. The pots are kept for 21 days at room temperature with 14 to 17 hours light (daylight or its equivalent) per day.

Determination of the herbicidal effect of the particular herbicide is made after the 21 day period. The determination involves a visual evaluation of the degree and quality of damage to the various seed plants.

The compounds of formula I are applied in the above manner at dosages equivalent to 1.4 and 5.6 kg of active agent/hectare.

Herbicidal activity, that is to say, significant damage to the test plants is observed.

Example 42: Weed control - Post-emergence treatment

A procedure similar to that employed in Example 41 is followed with the exception that the test compounds (herbicides) are applied when the plants are at the 2-4 leaf stage, the sowing of the plant seeds being staggered to ensure that the plants reach the 2-4 leaf stage at about the same time.

Again the compounds of formula I are applied in the above manner at dosages corresponding to 1.4 kg/ha and 5.6 kg/ha. The determination of the herbicidal effect is made 21 days after application of the test compounds and involves an analogous evaluation as described in Example 41. Herbicidal activity is observed.

Example 43

Representative compounds of the invention are evaluated in the following pre-emergence test procedure.

Seed dishes measuring 30 x 40 cm are filled to a depth of 6 cm with a mixture of peat culture substrate and sand. The exposed surface of the peat culture substrate and sand mixture is sprayed with an aqueous test liquid (e.g. formulated in accordance with Example B) comprising a compound of the invention in a given concentration. The spray volume corresponds to 600l aqueous test liquid/ha. The same test is repeated with various concentrations of test liquid, whereby the concentrations are selected in such a manner that the desired application rates are realized. Six species of seed are then sown in each dish. The number of seeds sown for each plant species depends on the seed germination potential and also the initial growth size of the particular seed plant. After sowing of the seeds, the treated surface is covered with a thin layer about 0.5 cm deep of the peat culture substrate and sand mixture.

The prepared seed dishes are kept for 28 days at a temperature of 20 to 24°C and 14 to 17 hours light each day.

Determination of the herbicidal effect of the particular compound of the invention is made after the 28 day period. The determination involves a visual evaluation of the degree and quality of damage to

the various plants. Particular advantageous herbicidal properties are i.a. observed with the Compound Nos. 25, 26, 47, 48, 55, 75, 84 and 86 of Table A.

Some of the results obtained with an application rate corresponding to 1 kg active ingredient/hectare are summarised in the following Table B.

Example 44: Post-emergence Treatment

A further evaluation of representative compounds of the formula I is effected in a post-emergence test procedure similar to that of the pre-emergence test described in Example 43, except that the herbicide test liquid is applied when the plants are at a 2-4 leaf

stage. For that purpose the various plants species are sown in time-staggered relationship. The greenhouse conditions (temperature, light) are as in Example 43. Determination of the herbicidal effect is also effected 28 days after application according to the method of Example 41.

Particular advantageous herbicidal properties are i.a. observed with the Compound Nos. 25, 26, 47, 48, 55, 75, 84 and 86 of Table A. Some of the results obtained with application rates corresponding to 5 kg active ingredient/hectare are summarised in the following Table C.

Plant treated	Pre-emergence application 1 kg/ha							
	Compound Tested - % damage							
	25	26	47	48	55	75	84	86
Amaran. retrofl.	90	90	100	100	100	100	80	99
Capsella b.p.	80	100	100	80	90	70	70	90
Chenop. alb.	80	80	90	50	20	50	60	60
Galium aparine	80	10	50	0	10	10	50	10
Senecio vulg.	90	80	80	80	80	100	90	100
Stellaria media	90	70	70	80	50	50	70	90
Alfalfa	80	60	60	50	80	10	70	90
Bean	0	0	20	20	0	0	10	10
Carrot	80	90	90	90	70	100	80	90
Cotton	20	0	10	0	0	0	20	0
Flax	80	30	60	50	30	0	20	10
Potato	0	10	10	0	0	0	0	0
Soya	50	10	30	30	0	10	30	30
Sugar beet	60	10	10	0	0	0	20	30
Rape	10	0	20	0	0	0	20	10
Sunflower	0	10	30	50	0	0	0	10
Agropyron repens	90	100	90	60	80	90	90	70
Agrostis alba	100	100	100	100	100	100	100	100
Alopec. myos.	90	90	80	20	50	80	80	90
Apera sp.venti.	100	100	100	100	100	100	100	100
Avena fatua	80	80	80	40	80	90	50	90
Echinochloa c.g.	90	100	90	90	90	100	90	90
Corn	20	30	50	30	0	50	90	60
Wheat	70	100	90	90	0	100	60	70

TABLE C Post-emergence application 5 kg/ha								
Plant treated	Compound Tested - % damage							
	25	26	47	48	55*	75	84	85*
Amaran. retrofl.	90	80	70	50	80	60	80	90
Capsella b.p.	80	60	20	20	80	50	80	60
Chenop. alb.	90	40	40	20	20	40	30	60
Galium aparine	80	60	40	30	80	20	70	60
Senecio vulg.	80	50	70	80	90	90	80	90
Stellaria media	90	50	60	10	30	40	60	40
Alfalfa	80	60	20	10	50	20	70	70
Bean	100	20	30	20	20	30	30	50
Carrot	70	90	30	100	80	100	100	80
Cotton	70	60	50	40	50	60	70	70
Flax	80	70	90	100	80	90	40	80
Potato	70	30	20	10	10	10	40	10
Soya	90	30	30	30	20	30	30	50
Sugar beet	30	20	70	0	10	10	0	10
Rape	40	20	10	10	30	60	50	40
Sunflower	60	50	30	80	40	60	90	50
Agropyron repens	70	30	70	50	50	50	30	50
Agrostis alba	-	-	-	-	-	-	-	-
Alopec. myos.	80	90	80	70	60	90	80	90
Apera sp.venti.	90	100	100	100	100	100	90	90
Avena fatua	90	100	90	80	90	100	100	100
Echinochloa c.g. 1)	80	60	80	80	90	70	80	90
2)	90	80	80	90	100	80	90	100
Corn	80	60	100	90	30	30	70	80
Wheat	70	90	80	50	10	60	60	80
Rice 2)	40	30	10	50	30	30	30	40

* 4 kg/ha 1) upland conditions; 2) paddy conditions

Field Test

Compound No. 25 has further been evaluated in a pre-emergence Multicrop-Fieldscreening under the following conditions:

- 5 - Crops Soybean ('Steele') - (So)
Cotton ('Stoneville 213') - (Cot)
Corn (LG 11) - (Co)
Wheat (Svenno) - (Wh)
- Weeds Alopecurus pratensis - (Al)
10 Echinochloa crus galli - (Ech)
Galium aparine
Chenopodium album and polyspermum } dicots
Amaranthus retroflexus
- (All the weeds, except for Alopecurus pratensis which was sown, were naturally occurring).
- Spray volume: 750 lt/ha
- Replicates: 3
- Evaluation: 28 days after application
- Standards: Alachlor (=α-chloro-2',6'-diethyl-N-methoxymethyl-acetanilide) and Metolachlor [=α-chloro-2'-ethyl-6'-methyl-N-(1-methyl-2-methoxyethyl)acetanilide].
- 20

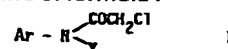
The following herbicidal activity was observed:

Compound	kg/ha	So	Cot	Co	Wh	Al/Ech	Dicots
No. 25	0.6	0	0	10	63	92	43
	1.25	3	3	53	80	100	67
	2.5	5	7	80	92	100	87
Alachlor	1.25	0	0	0	10	43	43
	2.5	0	3	0	53	78	57
Metolachlor	1.25	0	0	0	33	100	30
	2.5	0	0	0	73	100	43

- 25 The above results indicate an herbicidal activity which is equal or superior to that found with commercially available standards against monocotyledonous weeds, and is superior to that of such standards against dicotyledonous weeds. The herbicidal activity is selective in soybean, cotton and at the lowest application rate also in corn.

CLAIMS

1. A compound of formula I



- 35 Wherein Ar is a 5-membered heteroaromatic group comprising 1 or 2 ring heteroatoms selected from O, S and N, and linked by a C-ring atom to the N-atom of the N(Y)COCH₂Cl group to which it is bound, whereby, where Ar is pyrazolyl, said N(Y)COCH₂Cl group is in the 4-position, and Y is allene, CH₂-CH=C=CH₂.
- 40

or is a hydrocarbon selected from C₁₋₈alkyl, C₃₋₈alkenyl, C₃₋₈alkynyl, C₃₋₈cycloalkyl, C₆₋₈cycloalkenyl, C₃₋₈cycloalkyl - C₁₋₅alkyl, whereby such hydrocarbon is unsubstituted or substituted by

5 halogen selected from F, Cl or Br; or is a group CH(R₁)-COY₁,

wherein R₁ is H or C₁₋₅alkyl and

Y₁ forms together with the CO-group to which it is bound an ester or amide function;

10 or is a group R₂-Az,

wherein R₂ is CH₂ or CH₂-CH₂ unsubstituted or substituted by C₁₋₅alkyl and

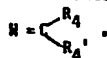
Az is an aromatic heteroring selected from a di- or triazole linked by one of its nitrogen atoms to R₂, a

15 5-membered heteroring linked by a C-atom of said ring to R₂ and having 1 to 3 heteroatoms selected from the group consisting of O, S or N, and a pyrimidine group, or is a 2-oxo-1-pyrrolidinyl group in which one CH₂ group can be replaced by O,

20 S or NCH₃ and 5-oxo- and/or bicyclic benz[c] fused derivatives of such 2-oxo-1-pyrrolidinyl group or is a group A-O-R₃,

wherein R₃ is H, hydrocarbon selected from the group consisting of C₁₋₅alkyl, C₃₋₈alkenyl,

25 C₃₋₈alkynyl, C₃₋₈cycloalkyl, C₆₋₈cycloalkenyl or C₃₋₈cycloalkyl - C₁₋₅alkyl, whereby the hydrocarbon is unsubstituted or substituted; or a group



30 wherein R₄ is hydrocarbon selected from C₁₋₅alkenyl, C₃₋₅alkynyl, C₃₋₈cycloalkyl, C₆₋₈cycloalkenyl, C₃₋₈cycloalkyl - C₁₋₅alkyl, whereby the hydrocarbon is unsubstituted or substituted by halogen selected from F, Cl or Br; or is allene

35 R₄' is H or has one of the meanings defined for R₄ and A is a hydrocarbon moiety, which may be linked with R₃ to form a saturated oxygen containing heterocyclic ring comprising 1 or 2 oxygens as heteroatom, and whereby the N and O atoms to

40 which it is bound are separated by up to 3 C-atoms; or is a group CH-C = NOR₄,



wherein R₄ is as defined above,

45 R₅ and R₅' are independently H or CH₃ or R₅ together with R₅' are (CH₂)₃ or (CH₂)₄

or is a group CH-N(CH₃)COCH₃



50 wherein R₆ is H or C₁₋₃alkyl.

2. A compound according to Claim 1, wherein Ar is at least substituted in *o*-position of the chloroacetamide group by a substituent selected from the group consisting of halogen; C₁₋₄alkyl unsubstituted

55 or substituted by halogen, C₁₋₄alkoxy or C₃₋₆cycloalkyl; C₂₋₄alkenyl unsubstituted or substituted by C₁₋₄alkoxy; C₃₋₆cycloalkyl; formyl or C₂₋₄alkanoyl; C(=NOC₁₋₄alkyl) - C₁₋₃alkyl; C(OC₁₋₄alkyl)₂ - C₁₋₃alkyl; CH(OC₁₋₄alkyl)₂;

60 C₁₋₄alkyl-S; C₁₋₄alkyl-SO; C₁₋₄alkyl-SO₂; C₁₋₄alkoxy-carbonyl; C₁₋₄alkoxy unsubstituted or substituted by halogen or C₁₋₄alkoxy; C₂₋₄alkenyloxy; C₂₋₄alkinyloxy; HO and CH₂OH and esters thereof.

65 3. A compound according to Claim 2, wherein Ar contains either one ring heteroatom or two adjacent ring heteroatoms, whereby the heteroaromatic ring is at least disubstituted in *o,o'*-position of the chloroacetamide group by substituents selected from the group as defined in Claim 2, any optional

70 additional substituent of said heteroaromatic ring being selected from C₁₋₄alkyl, halogen and C₁₋₄alkoxycarbonyl.

4. A compound according to Claim 3, wherein Ar is a substituted thiophene, isothiazole or isoxazole ring.

5. A compound according to Claim 4, wherein Ar is a substituted thiophene or isothiazole ring.

6. A compound according to Claim 5, wherein Ar is substituted thiophene.

7. A compound according to Claim 6, wherein Ar is 3-thienyl 2,4-disubstituted by substituents selected from C₁₋₄alkyl and C₁₋₄alkoxy.

8. A compound according to any one of Claims 1 to 7 wherein Y is a group R₂-Az or a group A-OR₃ (R₂, Az, A and R₃ being as defined in Claim 1).

9. A compound according to Claim 8, wherein R₂ is CH₂ or CH(CH₃), Az is 1-pyrazolyl or 3,5-diCH₃-1-pyrazolyl, A is CH₂, CH₂-CH₂ or a monomethylated derivative thereof and R₃ is C₁₋₃alkyl.

10. A compound of formula I according to Claim 9, selected from the group of compounds wherein Ar and Y are

a) 2,4-dimethyl-thien-3-yl and pyrazolyl-1-

95 methyl resp.

b) 2,4-dimethyl-thien-3-yl and 1-(pyrazolyl-1) ethyl resp.

c) 2,4-dimethyl-thien-3-yl and CH₂OC₂H₅ resp.

d) 2,4-dimethyl-thien-3-yl and CH₂OC₃H₇ resp.

100 e) 2,4-dimethyl-thien-3-yl and CH(CH₃)CH₂OCH₃ resp.

f) 2-methyl-4-ethyl-thien-3-yl and CH₂OC₂H₅ resp.

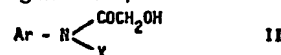
g) 2-methyl-4-methoxy-thien-3-yl and

105 pyrazolyl-1-methyl resp.

h) 2-methyl-4-methoxy-thien-3-yl and CH₂CH₂OC₂H₅ resp.

11. Process for the production of a compound of formula I stated in Claim 1, comprising

110 a) substituting in a compound of formula II



wherein Ar and Y are as defined above,

the HO group of the N-hydroxyacetyl group by Cl,

115 b) reacting a compound of formula III



wherein Ar is as defined above,

with a compound of formula IV

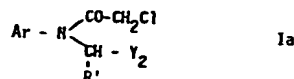


wherein Y is as defined above,

and L is a leaving group capable of being split off

under the N-alkylation reaction conditions,

c) obtaining a compound of formula Ia

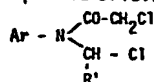


wherein R' is H or C₁₋₃alkyl

Y₂ is Az', OR₃, N(CH₃)COCH₃ and

Az' is a di- or triazole linked by one of its nitrogen atoms to CHR'

- 5 and Ar and R₃ are as defined above,
by reacting a compound of formula V



- 10 wherein Ar and R' are as defined above,
with a reactive derivative of a compound of formula VI



wherein Y₂ is as defined above,

- 15 d) N-acylation of a compound of formula VII



wherein Ar and Y are as defined above,
with chloroacetyl chloride, or a reactive functional derivative thereof.

- 20 12. A compound as stated in Claim 1, whenever
obtained by a process claimed in Claim 11.

13. The use of a compound of any one of Claims
1 to 10 and 12 as herbicide.

14. The use of a compound of any one of Claims

- 25 1 to 10 and 12 as selective herbicide.

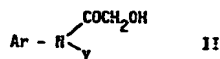
15. The use according to Claim 14 in sugar beets,
cotton, potatoes, soybeans, corn or peanuts.

16. A herbicidal composition comprising a com-
pound of any one of Claims 1 to 10 and 12 in
association with an herbicidally acceptable diluent.

- 30 17. The use according to any one of Claims 13 to
15 substantially as described herein.

18. A composition according to Claim 16 sub-
stantially as described herein.

- 35 19. A compound of formula II



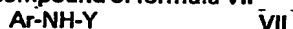
wherein Ar and Y are as defined in Claim 1.

- 40 20. A compound of formula III



wherein Ar is as defined in Claim 1.

21. A compound of formula VII



- 45 wherein Ar and Y are as defined in Claim 1.

22. A 3-aminothiophene substituted in 2- and
4-position by a group selected from C₁₋₄alkyl and
C₁₋₄alkoxy.

**This Page is Inserted by IFW Indexing and Scanning
Operations and is not part of the Official Record**

BEST AVAILABLE IMAGES

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images include but are not limited to the items checked:

- ☐ **BLACK BORDERS**
- ☐ **IMAGE CUT OFF AT TOP, BOTTOM OR SIDES**
- ☐ **FADED TEXT OR DRAWING**
- ☐ **BLURRED OR ILLEGIBLE TEXT OR DRAWING**
- ☐ **SKEWED/SLANTED IMAGES**
- ☐ **COLOR OR BLACK AND WHITE PHOTOGRAPHS**
- ☐ **GRAY SCALE DOCUMENTS**
- ☒ **LINES OR MARKS ON ORIGINAL DOCUMENT**
- ☐ **REFERENCE(S) OR EXHIBIT(S) SUBMITTED ARE POOR QUALITY**
- ☐ **OTHER:** _____

IMAGES ARE BEST AVAILABLE COPY.

As rescanning these documents will not correct the image problems checked, please do not report these problems to the IFW Image Problem Mailbox.